Supplemental Table S1. Strains and plasmids used in this study

Name	Description	Source or Note
Strains		
DH10b	E. coli K12 F $^ \lambda^-$ mcrA Δ (mrr-hsdRMS-mcrBC) $\phi 80\Delta lacZM15$ $\Delta lacX74$ recA1 araD139 Δ (araA-leu)7697 galE15 galK16 rpsL endA1 nupG	(1)
BL21λDE3	E. coli B F ompT gal dcm lon hsdS(rB mB λDE3(lacUV5::T7 gene 1 imm21 int Δnin5 ΔEcoRI[21226-26104] BamHI27972)	(2)
<u>Plasmids</u>		
pRARE2	Cm ^R Rosetta plasmid encoding rare tRNAs	Novagen
pET28a	Kn ^R ColEI <i>ori</i> encodes <i>E. coli lacI</i> , T7 promoter, and polylinker	Novagen
pAC22	pET28a derivative. expresses <i>M. bovis</i> ω , α , and β :: β' fusion with C-terminal His ₈ tag from T7 promoter.	
pRM629	pET28a derivative; expresses proteins with N-terminal His ₁₀ -ppx-tag from T7 promoter; Kn ^R . (ppx, Precission protease site)	
pAC27	pRM629 with <i>M. bovis</i> σ^{A} with N-terminal His ₁₀ ppx tag	
pVS9	expresses E. coli EcoRIQ111 from T7 pET33 based promoter	I. Artsimovitch
pAC81	pRM629 with <i>M. bovis</i> NusA with N-terminal His ₁₀ ppx tag	
pAC82	pRM629 with <i>M. bovis</i> NusG N-terminal His ₁₀ ppx tag	
pAC83	pRM629 M. bovis NusG ΔMboi1	
pAC84	pRM629 M. bovis NusG Δ Mboi2	
pAC85	pRM629 M. bovis NusG ΔCTD	
pRLG3748	p770 with -35 consensus promoter followed by <i>rrnB</i> terminators T1 and T2	(3)
pAC47	pRLG3748 with introduced BamHI and NcoI sites before and after rrnBT1 terminator respectively	
pAC48	pAC47 with M. tuberculosis tuf terminator	
pAC49	pAC47 with M. tuberculosis Rv1324 terminator	
pAC50	pAC47 with M. smegmatis gyrA terminator	
pAC51	pAC47 with E. coli trpL terminator	
pAC52	pAC47 with M. tuberculosis rpoC terminator	
pAC53	pAC47 with E. coli rpoC terminator	
pAC54	pAC47 with M. tuberculosis Rv3444c terminator	
pAC55	pAC47 with M. bovis fadD23 terminator	
pAC56	pAC47 with <i>M. smegmatis gyrA*</i> sequence from Ref. 4.	(4)
pAC58	pAC47 with M. bovis asnB terminator	
pAC59	pAC47 with <i>M. bovis asnB</i> -1 terminator	
pAC60	pAC47 with <i>M. bovis asnB</i> -2 terminator	
pAC61	pAC47 with M. bovis asnB-3 terminator	
pAC62	pAC47 with <i>M. bovis asnB</i> -4 terminator	
pAC63	pAC47 with <i>M. bovis asnB</i> -5 terminator	

pAC64	pRLG374 with one EcoRI before promoter changed to PvuI and other EcoRI introduced between promoter and terminator
pAC65	pAC51 with one EcoRI before promoter changed to PvuI and other EcoRI introduced between promoter and terminator
pAC66	pAC48 with one EcoRI before promoter changed to PvuI and other EcoRI introduced between promoter and terminator
pAC67	pAC49 with one EcoRI before promoter changed to PvuI and other EcoRI introduced between promoter and terminator
pAC68	pAC54 with one EcoRI before promoter changed to PvuI and other EcoRI introduced between promoter and terminator
pAC69	pAC52 with one EcoRI before promoter changed to PvuI and other EcoRI introduced between promoter and terminator
pAC70	pAC48 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC71	pAC49 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC72	pAC52 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC73	pAC50 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC74	pAC56 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC75	pAC58 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC76	pAC59 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC77	pAC60 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC78	pAC61 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC79	pAC62 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC80	pAC63 but C at positions +11,14,16, and 20 changed to G, T, G, and G respectively
pAC87	pAC70 but with t _{synA} terminator
pAC88	pAC70 but with t _{synB} terminator

- 1. **Grant SG, Jessee J, Bloom FR, Hanahan D.** 1990. Differential plasmid rescue from transgenic mouse DNAs into Escherichia coli methylation-restriction mutants. Proc. Natl. Acad. Sci. USA **87:**4645-4649.
- 2. **Studier FW, Moffatt BA.** 1986. Use of bacteriophage T7 RNA polymerase to direct selective high-level expression of cloned genes. J. Mol. Biol. **189:**113-130.
- 3. Gaal T, Ross W, Estrem ST, Nguyen LH, Burgess RR, Gourse RL. 2001. Promoter recognition and discrimination by EsigmaS RNA polymerase. Mol. Micro. 42:939-954.
- 4. **Unniraman S, Prakash R, Nagaraja V.** 2001. Alternate paradigm for intrinsic transcription termination in eubacteria. J. Biol. Chem. **276:**41850-41855.